

(FILE 'HOME' ENTERED AT 14:45:05 ON 27 JAN 2003)

FILE 'MEDLINE' ENTERED AT 14:45:19 ON 27 JAN 2003

L1 0 S SHAW G?/AU AND ADAPTER  
L2 835 S SHAW G?/AU  
L3 2 S L2 AND SH2  
L4 19 S L2 AND BIND  
L5 8 S L2 AND PLECKSTRIN  
L6 513 S PLECKSTRIN HOMOLOGY DOMAIN  
L7 11 S L6 AND (MICROSPHERE OR MICROPARTICLE OR NANOSPHERE  
OR NANOPAR

FILE 'CAPLUS, EMBASE, BIOSIS, BIOTECHDS, SCISEARCH' ENTERED AT  
14:52:07

ON 27 JAN 2003

L8 24 S L7  
L9 17 DUP REM L8 (7 DUPLICATES REMOVED)  
L10 30 S (MICROSPHERE OR MICROPARTICLE OR NANOSPHERE OR  
NANOPARTICLE O  
L11 21 DUP REM L10 (9 DUPLICATES REMOVED)  
L12 4 S L11 AND TARGET?

FILE 'MEDLINE' ENTERED AT 15:05:16 ON 27 JAN 2003

L13 20150 S (MICROSPHERE OR MICROPARTICLE OR NANOPARTICLE OR  
NANOSPHERE)  
L14 44590 S L13 OR LIPOSOME  
L15 3138 S L14 AND TARGET?  
L16 3 S L15 AND ADAPTER

FILE 'CAPLUS, EMBASE, BIOSIS, BIOTECHDS, SCISEARCH' ENTERED AT  
15:08:40

ON 27 JAN 2003

L17 13 S L16  
L18 11 DUP REM L17 (2 DUPLICATES REMOVED)

=> file stnguide

COST IN U.S. DOLLARS

(FILE 'HOME' ENTERED AT 12:09:15 ON 30 JAN 2003)

FILE 'MEDLINE' ENTERED AT 12:09:22 ON 30 JAN 2003

L1 68 S RIBONUCLEASE I  
L2 1 S L1 AND MOUSE

FILE 'STNGUIDE' ENTERED AT 12:13:56 ON 30 JAN 2003

FILE 'STNGUIDE' ENTERED AT 12:27:31 ON 30 JAN 2003

FILE 'MEDLINE' ENTERED AT 12:27:38 ON 30 JAN 2003

L3 1 S HUMAN RIBONUCLEASE A  
L4 0 S HUMAN RIBONUCLEASE I  
L5 0 S HUMAN RIBONUCLEASE I  
L6 0 S RIBONUCLEASE A AND RIBONUCLEASE I  
L7 4 S RIBONUCLEASE A AND RIBONUCLEASE I  
L8 64 S RAINES R7/AU AND RIBONUCLEASE  
L9 0 S L8 AND RNASE A AND RNASE I

FILE 'CAPLUS, EMBASE, BIOSIS, BIOTECHDS, SCISEARCH' ENTERED AT  
12:32:00

ON 30 JAN 2003

L10 5 S L9  
L11 2 DUP REM L10 (3 DUPLICATES REMOVED)

(FILE 'HOME' ENTERED AT 13:32:21 ON 30 JAN 2003)

FILE 'MEDLINE' ENTERED AT 13:32:31 ON 30 JAN 2003

L1 20 S VITRONECTIN AND (VASCULAR ENDOTHELIAL CELL OR  
VASCULAR ENDOTHE  
L2 741 S VITRONECTIN AND TUMOR  
L3 13 S L2 AND SOLID TUMOR

FILE 'STNGUIDE' ENTERED AT 13:49:04 ON 30 JAN 2003

=> d his

(FILE 'HOME' ENTERED AT 11:03:06 ON 31 JAN 2003)

FILE 'MEDLINE' ENTERED AT 11:03:13 ON 31 JAN 2003

L1 128 S BOVINE RIBONUCLEASE OR BOVINE RNASE  
L2 0 S BOVINE RIBONUCLEASE I OR BOVINE RNASE I  
L3 4 S BOVINE (5A) RIBONUCLEASE I OR BOVINE (5A) RNASE I

S #

Updt

Database

Query

Time

Comment

S5938

U  
USPT

4885172.pn. and  
(excipient or carrier)  
2003-01-31  
09:15:34

S5937  
U  
USPT

(ligand same fusion  
protein same target\$ )  
and targeting ligand  
2003-01-30  
14:11:57

S5936  
U  
USPT

ligand same fusion  
protein same target\$  
2003-01-30  
14:11:23

S5935  
U  
USPT

ligand same fusion  
protein  
2003-01-30  
14:11:04

S5934  
U  
USPT

lignad same fusion  
protein  
2003-01-30  
14:10:51

S5933  
U  
USPT

targeting ligand same  
fusion  
2003-01-30  
14:06:21

S5932  
U  
USPT

targeting ligand same

fusion protein  
2003-01-30  
14:01:12

S5931  
U  
USPT

pretargeting and S  
protein  
2003-01-30  
11:04:45

S5930  
U  
USPT

6015897.pn. and  
targeting moiety\$  
2003-01-30  
10:57:41

S5929  
U  
USPT

6015897.pn. and (s  
protein or s peptide)  
2003-01-30  
10:56:39

S5928  
U  
USPT

targeting moiety and  
vegf and ( targeting  
moiety same (vegf or  
vascular endothelial))  
2003-01-30  
10:52:11

S5927  
U  
USPT

targeting moiety and  
(vegf or vascular  
endothelial) and (  
targeting moiety same  
(vegf or vascular  
endothelial))  
2003-01-30  
10:50:28

S5926  
U

USPT

targeting moiety same  
(vegf or vascular  
endothelial)

2003-01-30

10:48:23

S5925

U

PGPB,JPAB,EPAB,DWPI,TDBD

targeting moiety same  
(vegf or vascular  
endothelial)

2003-01-30

10:48:13

S5924

U

PGPB,JPAB,EPAB,DWPI,TDBD

targetting moiety same  
(vegf or vascular  
endothelial)

2003-01-30

10:47:58

S5923

U

PGPB,JPAB,EPAB,DWPI,TDBD

(tropism same vascular  
endothelial )

2003-01-30

10:42:12

S5922

U

USPT

tropism same vascular  
endothelial

2003-01-30

10:41:43

S5921

U

USPT

tropism same vegf

2003-01-30

10:41:18

S5920

U

PGPB,JPAB,EPAB,DWPI,TDBD

fiber protein and knob

and vegf

2003-01-30

10:37:06

S5919

U

USPT

fiber protein and knob

and vegf

2003-01-30

10:29:09

L1 ANSWER 61 OF 68 MEDLINE  
AN 64032753 MEDLINE  
DN 64032753  
TI STUDIES ON B. SUBTILIS **RIBONUCLEASE. I.**  
CHARACTERIZATION OF ENZYMATIC SPECIFICITY.  
AU RUSHIZKY G W; GRECO A E; HARTLEYRW J R; SOBER H A  
SO BIOCHEMISTRY, (1963 JUL-AUG) 128 787-93.  
ISSN: 0006-2960.  
CY United States  
DT Journal  
LA English  
FS OLDMEDLINE  
EM 196403  
ED Entered STN: 19990716  
Last Updated on STN: 19990716

L1 ANSWER 22 OF 68 MEDLINE  
 AN 77140877 MEDLINE  
 DN 77140877 PubMed ID: 321440  
 TI Effects of polyamines on the activities of Escherichia coli  
**ribonuclease I** and II.  
 AU Kumagai H; Igarashi K; Yoshikawa M; Hirose S  
 SO JOURNAL OF BIOCHEMISTRY, (1977 Feb) 81 (2) 381-8.  
 Journal code: 0376600. ISSN: 0021-924X.  
 CY Japan  
 DT Journal; Article; (JOURNAL ARTICLE)  
 LA English  
 FS Priority Journals  
 EM 197705  
 ED Entered STN: 19900313  
 Last Updated on STN: 19970203  
 Entered Medline: 19770527  
 AB The effects of polyamines on the breakdown of synthetic polynucleotides  
 [poly(A), poly(C), and poly(U)] by E. coli **ribonuclease**  
**I** [ribonuclease 3'-oligonucleotidohydrolase, EC 3.1.4.23] and  
 ribonuclease II [EC 3.1.4.1] have been studied. The degradation of poly(C)  
 by RNase II was stimulated by spermine and spermidine, while that of  
 poly(A) by RNase II was not affected by polyamines. Under our standard  
 experimental conditions, the breakdown of poly(U) by RNase II was  
 inhibited slightly by polyamines. The stimulatory effect of spermine and  
 spermidine on the breakdown of poly(C) occurred in the absence of  
 monovalent cations but not in the absence of divalent cations. When  
 polyamines were used as a stimulant of RNase II, the ratio of poly(C)  
 degradation to poly(U) degradation was greater in the presence of  
 inhibitors such as poly(G) than in their absence. Although the breakdown  
 of all synthetic polynucleotides by RNase I was stimulated by polyamines,  
 the degree of stimulation by polyamines was in the order poly(C) greater  
 than poly(A) (see text) poly(U). However, the difference in degree of  
 stimulation among polynucleotides decreased as monovalent cation  
 concentration was increased.



L1 ANSWER 21 OF 68 MEDLINE  
 AN 77166629 MEDLINE  
 DN 77166629 PubMed ID: 16063  
 TI Epidermal nucleases. II. The multiplicity of ribonucleases in guinea-pig epidermis.  
 AU Melbye S W; Freedberg I M  
 SO JOURNAL OF INVESTIGATIVE DERMATOLOGY, (1977 May) 68 (5) 285-92.  
 Journal code: 0426720. ISSN: 0022-202X.  
 CY United States  
 DT Journal; Article; (JOURNAL ARTICLE)  
 LA English  
 FS Abridged Index Medicus Journals; Priority Journals  
 EM 197706  
 ED Entered STN: 19900313  
 Last Updated on STN: 19950206  
 Entered Medline: 19770630  
 AB Ribonuclease activity has been extracted from adult guinea-pig epidermis by sequential homogenization in dilute sodium acetate and sulfuric acid. The extracts were subjected to ammonium sulfate fractionation and to affinity and ion exchange chromatography. Three **ribonucleases** (I, II, III) were separated from the sodium acetate extract and 6(A, B1, B2, B3, C, D) were isolated from the sulfuric acid extract. The degree of purification varies from 65-fold to 8,700-fold and the apparent molecular weights of the active forms of 8 of the 9 ribonucleases range from 10,000 to 36,500. No phosphodiesterase activity is present in any of the 9 fractions, but there is alkaline phosphatase activity in one (I) and deoxyribonuclease activity in a second (B3). Two of the ribonucleases have acid pH optima (a1, B3), while the others are most active between PHs 6.8 and 7.8. The activity of 4 of the fractions is sensitive to added EDTA (III, A, B2, B3), but no stimulatory metal ions were found. Low concentrations of the polyamine spermidine enhanced the activity of 3-fractions (III, C, D). Yeast ribonucleic acid is degraded exonucleolytically by 2 fractions (I, A) and endonucleolytically by the remaining 7. In experiments with homopolyribonucleotide substrates, poly U was generally the preferred substrate. Substantial hydrolysis of poly A occurred with 2 fractions (A, B3) and slight hydrolysis of poly G with 2 other fractions (B2, C).

L3 ANSWER 5 OF 6 BIOTECHDS COPYRIGHT 2003 THOMSON DERWENT AND ISI  
AN 1996-02111 BIOTECHDS  
TI Strategies to accomplish targeted gene delivery employing  
tropism-modified recombinant adeno viral vectors;  
adeno virus vector construction for tissue-specific gene expression  
and gene therapy (conference abstract)  
AU Michael S I; Douglas J T; Miller C R; Krasnykh V; Hong J S; Engler J A;  
Curriel D T  
CS Univ.Alabama  
LO Department of Biochemistry and Molecular Genetics, University of Alabama  
at Birmingham, Birmingham, AL 35294, USA.  
SO Cancer Gene Ther.; (1995) 2, 4, 321  
CODEN: 2815V ISSN: 0929-1903  
Gene Therapy of Cancer, 4th International Conference, San Diego, CA, USA,  
9-11 November, 1995.  
DT Journal  
LA English  
AB Targeting ligands were introduced within an adeno virus (AV) fiber  
protein, which mediates target cell receptor binding. Fiber-ligand  
fusions which retained the native fiber quaternary structure, accumulated  
in HeLa cell nuclei and presented the targeting ligand on the fiber  
ecto-domain. Fusions were incorporated into AV vectors by homologous  
recombination, and transient expression was used to pseudotype the fiber  
protein in vitro. Anti-fiber antibody (Ab)-ligand fusion proteins were  
used to re-target the virus. Anti-fiber **knob** antibodies (Abs)  
bound with high affinity to the fiber **knob** domain and ablated  
AV binding to endogenous target cell receptors. Single chain variable  
regions of these Abs were fused with a specific ligand domain to form  
bispecific Abs which blocked endogenous AV binding and had novel  
specificity. A ligand binding domain was attached to AV capsid proteins  
via a **biotin-streptavidin** bridge. AV capsid proteins  
could be extensively biotinylated without impairing binding or reporter  
gene expression. These AV vectors should allow targeted cell-specific  
delivery in vivo. (0 ref)

For Response to 11.2/2 RET

L3 ANSWER 3 OF 4 MEDLINE  
AN 76018943 MEDLINE  
DN 76018943 PubMed ID: 240382  
TI Correlation proton magnetic resonance studies at 250 MHz of **bovine** pancreatic **ribonuclease**. I. Reinvestigation of the histidine peak assignments.  
AU Markley J L  
SO BIOCHEMISTRY, (1975 Aug 12) 14 (16) 3546-54.  
Journal code: 0370623. ISSN: 0006-2960.  
CY United States  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
EM 197512  
ED Entered STN: 19900313  
Last Updated on STN: 19970203  
Entered Medline: 19751211  
AB The deuterium exchange kinetics of the C(2) protons of the four histidine residues of native bovine pancreatic ribonuclease A have been followed at pH 6.5 and 8.0 by proton magnetic resonance spectroscopy (1H NMR). Comparison of the order of exchange of the histidine peaks with tritium exchange rates into individual histidine residues [Ohe, M., Matsuo, H., Sakiyama, F., and Narita, K. (1974), J. Biochem. (Tokyo) 75, 1197] supports the previous assignment of histidine NMR peaks H(1) and H(4) to histidine-105 and histidine-48 but requires reassignment of peaks H(2) and H(3) to histidine-119 and histidine-12, respectively. Ribonuclease A samples having differentially deuterated histidines have been used to verify the existence of crossover points in the histidine proton magnetic resonance titration curves and to observe the discontinuous titration curve of histidine-48. Proton magnetic resonance peaks have been assigned to the C(4) protons of the four histidine residues of ribonuclease A on the basis of their unit proton areas and by matching their titration shifts with the more readily visible C(2)-H peaks of the histidines. The pK' values derived from the C(4)-H data agree, within experimental limits, with those derived from C(2)-H data. The C(4)-H peaks were assigned to histidine-12, -48, -105, and -119 of ribonuclease A on the basis of their pH dependence, pK' values, shifts of their pK' values in the presence of inhibitor cytidine 3'-phosphate, and by comparison with the assignments of the histidine C(2)-H peaks above.

L3 ANSWER 4 OF 4 MEDLINE  
AN 69260123 MEDLINE  
DN 69260123 PubMed ID: 5801478  
TI Heavy atom-labelled derivatives of **bovine** pancreatic **ribonuclease**. I. Specific reactions of ribonuclease with N-acetylhomocysteine thiolactone and silver ion.  
AU Shall S; Barnard E A  
SO JOURNAL OF MOLECULAR BIOLOGY, (1969 Apr) 41 (2) 237-51.  
Journal code: 2985088R. ISSN: 0022-2836.  
Report No.: NASA-69260123.  
CY ENGLAND: United Kingdom  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals; Space Life Sciences  
EM 196909  
ED Entered STN: 19900101  
Last Updated on STN: 19970203  
Entered Medline: 19690930